



**ACTA**

**ACTA MORPHOLOGICA  
ACADEMIAE SCIENTIARUM HUNGARICAE**

**TOMUS I.**

**FASCICULUS 4.**

Patho-histological Laboratory (leader: A. Keri) of the State Institute for Radium and  
Roentgen Institute in Budapest (Director: Prof. Bela Wald)

## ALKALINE PHOSPHATASE ACTIVITY CHANGES IN ORGANS OF RATS KILLED WITH ROENTGEN-RAYS\*

Nándy Ró, Livia Ró and Vilmos Farkas

(Received: 4. Jan. 1951.)

In our Institute the effect on living tissues of gamma rays originating from the disintegration of radium and that of Roentgen rays has been subject of general studies. In the first place we have been investigating changes in the enzyme systems of irradiated animals and human tumors. In this paper we wish to report on the behavior of the alkaline phosphatase reaction performed on tissues of rats.

The phosphatases are dephosphorylising enzymes which split off  $PO_4$  from phosphoric acid esters by means of hydrolysis. Phosphatase can be found everywhere in vegetable and animal tissues. The phosphatases have been classified differently by different authors. In general 1. phosphomonoesterases, 2. phosphodiesterases, 3. pyrophosphatases and 4. apyrases are distinguished.

For phosphomonoesterases, based on how they react to alpha and beta glycerophosphate as substrate, what reaction they give to Mg and how they are distributed in different organs, Folley and Kay have put forward the following classification: Type  $A_1$  (alkaline phosphatase), optimal activity at pH 9. Type  $A_2$  (acid phosphatase), optimal activity ranging from pH 4.5 to pH 5. Type  $A_3$  (vegetable phosphatase), optimal activity at pH 6 and finally type  $A_4$  (acid erythrocyte phosphatase), optimal activity at about pH 6.

Gömöri - and independently from him Takumatsu - were the first to succeed in demonstrating histochemically acid and alkaline phosphatase.

In this paper we wish to report on those alkaline phosphatase investigations which we performed according to Gömöri's method - on organs of rats killed with lethal doses of rays. The reason why we selected rats for our first experiments was that the rat, contrary to other experimental animals, is capable of synthesising vitamin C. Irradiated animals namely lose a very great quantity of ascorbic acid through excretion, and the resulting vitamin C deficiency - especially in guinea pigs - leads to a very marked decrease of

\* Presented at the 16th Dec. 1950 meeting of Hungarian Pathologists.

phosphatase activity of tissues. For our experiments we used 20 couples of rats, mostly males. Their average weight was 150 g. and they were 8 to 10 months old. The rats were given a total irradiation of 2000 to 4000 r. in one lethal dose or in fractionated doses. For irradiation we used a Siemens Stabilivolt apparatus with Grounacher switch and with a Siemens-Doglas inlay tube. 180 kV, 15 mA irradiations were given, from a distance of 50 cm., using a 0.5 Cu filter. To each of the irradiated animals belonged a control rat, similar in bodily condition and kept under the same circumstances. The rats kept on

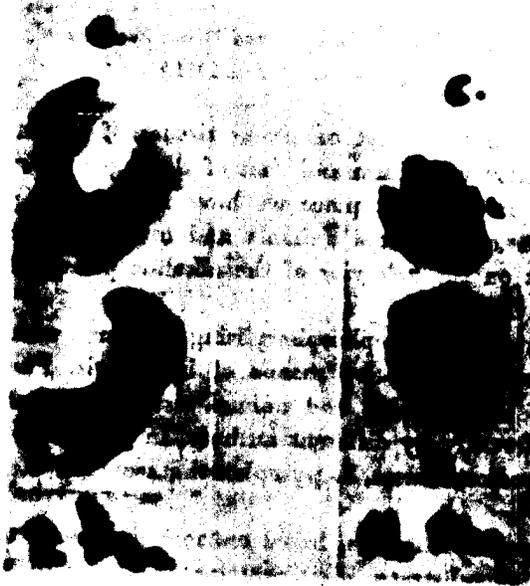


Fig. 1.  
Microscopic photographs of histological sections. c marks sections of the organ from control, & from experimental animals. Sections of the liver, after having performed the phosphatase reaction.

standard diet died generally after a period of 3 to 4 days following the irradiation. At 25 to 26 hours following the irradiation there was a marked loss of appetite, but the animals still kept on eating. When the irradiated animal died spontaneously, the control animal was also killed by a blow to the neckal region. Identical organs of both animals were embedded at the same time, the histological sections were mounted on the same slide and were treated with Gomori's method. For fixation 95 per cent ice-cooled alcohol, for embedding (quaternary) eups of alcohol, benzol and paraffine were used. Azro substrates contain glycerophosphate, as buffer sodium diethylbarbiturate were used. The precipitated  $\text{Ca}_3(\text{PO}_4)_2$  was made visible by means of Kossel's reaction.

PHOSPHATASE ACTIVITY IN THE LIVER OF RATS

In the histological sections in this way prepared we observed in irradiated animals a very marked increase of phosphatase activity. Most conspicuous was the greatly increased phosphatase activity of the liver, which could be detected even macroscopically in the sections (Fig. 1). The liver of the normal animal is according to Gombri's description and our observations made in the controls negative as regards phosphatase activity; in some cases there may be some activity limited to the biliary capillaries and to the walls of the vessels. (Fig. 2.) In irradiated animals we found very great phosphatase activity in the liver



Fig. 2.

The liver of a control. The perinuclear zone of phosphatase activity; phosphatase activity is seen in the wall of bile ducts and of greater vessels. (Same microscope, objective & ocular as Fig. 1.)

shows, showing nodular arrangement (Fig. 3). It was in the perinuclear zone of the liver cells and in the bile ducts surrounded by these cells that high degree of enzyme activity could be observed (Fig. 4). In the Kupffer cells, on the other hand, no phosphatase activity could be demonstrated. In other organs higher enzyme activity could be demonstrated in the membrane basalis of the testis, as well as in the bronchial epithel, in the edematous alveolar fluid and in the degenerated alveolar cells of the lungs. In the kidneys the results were more difficult to evaluate, since normal kidneys also show high-degree phosphatase activity. In the intestines no evaluations could be made, due to the rapid intestinal autolysis occurring in irradiated animals. Unfortunately we could not investigate the bone marrow. This and the central nervous system will be studied separately.

Summing up we may state that in the organs of rats killed with a lethal dose of Roentgen-rays there is an increase of the phosphatase activity. This increase is very marked in the liver.

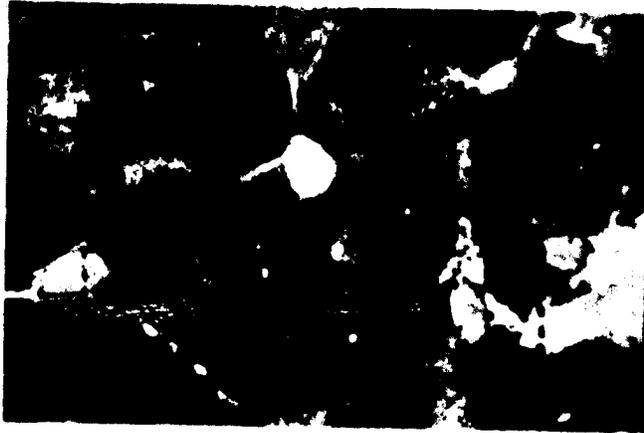


Fig. 3

General view of the liver of an irradiated animal. Zonal arrangement of phosphatase activity. The area in the square is shown in greater magnification in the next Fig. (Objective O, ocular 5)



Fig. 6

Greater magnification from Fig. 3 (Objective O, ocular 5 of a Zeiss microscope)

The occurrence of phosphatase activity in the liver may offer a partial explanation to the hypoglycemic conditions observed by Straus and Rother, Rother, and other authors, as well as to the disappearance of the liver glycogen following irradiation (Rother) in experimental animals.

The question arises: Why does the phosphatase activity of different organs—and mainly that of the liver—increase following irradiation? To the answer we must know the following data of literature:

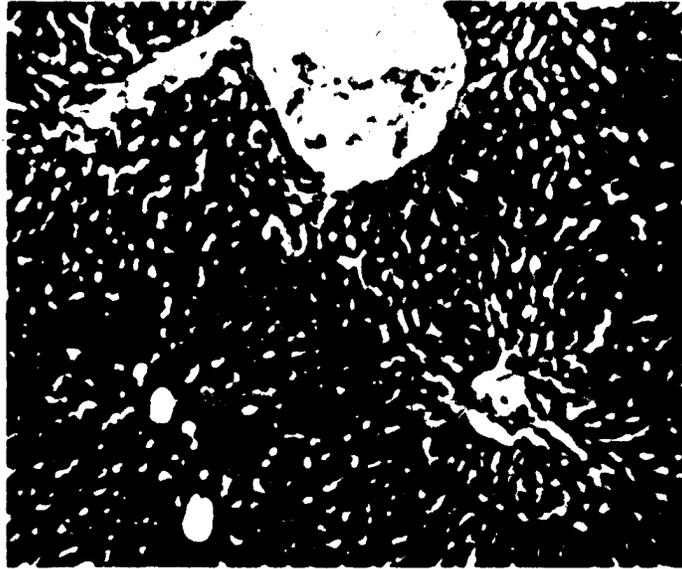


Fig. 1  
Higher magnification (Zeiss optics, objective 70x, ocular 30)

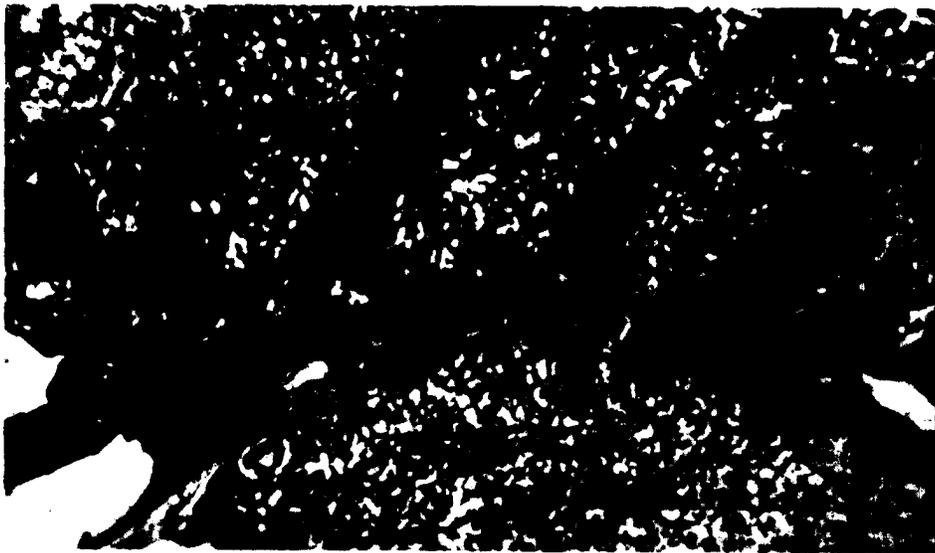


Fig. 2  
Microscopic view of the lung of an irradiated animal.

As a result of the irradiation, the activity of the enzyme is markedly reduced, the otherwise very high activity in the nucleohiston, this is probably due to depolymerisation. In our opinion, a similar process occurs in living cells on irradiation.

Holme, experimented with Jensen's rat sarcoma. He injected  $^{32}\text{P}$  isotope into the animal and found that on irradiation the nuclear metabolism, i.e. the incorporation of  $^{32}\text{P}$  into the thymonucleohiston, was materially reduced. Thus he confirmed Hevesy's previous observations made on animals having two tumors. Isolated irradiation of one of these tumors similarly decreased  $^{32}\text{P}$  uptake in the non-irradiated tumor.

Caspersson and Thorell have found that in the course of embryonal development the changes in phosphatase activity run parallel with the changes in concentration of nucleic acids respectively of nucleoproteids. It appears very likely that this enzyme takes part in the synthesis and destruction of nucleic acids.

Krugelis, Danelli and Catcheside have found in the cells of *Drosophila*, in the chromosomes, phosphatase activity in such a trabecular distribution which more or less corresponds to the arrangement of the Feulgen-positive bundles.

Finally it should be mentioned that in contrast to the former static theory also in the enzymology it is the dynamic conception that prevails, i.e. we consider enzymes to be effects and not separate substances. The same substance that in a certain state of the cell plays the part of a metabolite and is fuel supplying energy, under other circumstances may act as an enzyme. Our experiments and the quoted data of literature suggest that *the phosphatase activity occurring in the liver may be brought into connection with the depolymerisation of thymonucleohiston brought about by irradiation, which presents itself in the form of an enzyme effect.*

Further histochemical and other investigations are in progress with which we wish to obtain further evidence concerning the validity of this theory.

*Acknowledgements.* Authors express their thanks to K. Haffner, I. Dombi and Zs. Huppert, histotechnicians, for the valuable assistance.

#### Summary

In our Institute we investigate the effects of Roentgen and gamma rays on living tissues and on tumors. In the first place it was the changes of enzymes that were subjected to investigation by means of known histochemical reactions. In the present paper we report on the changes of alkaline phosphatase activity in tissues of rats killed with lethal doses of rays.

In our experiments rats were used, because these animals are capable of synthesizing vitamin C, in other animals irradiation with Roentgen rays gives rise to excessive vitamin C excretion and this reduces the phosphatase activity of their organs. The rats were irradiated with 3000-4000 r, in one dose or in fractionated doses (180 kV, 18 mA, 0.5 Cu, 50 cm). To each irradiated animal there was a control one being of the same bodily condition and kept under similar circumstances. When the irradiated animal died spontaneously, the control was also killed by a blow to the nuchal region. The organs of the two animals were prepared identically and at the same time. Identical organs were examined on one slide. The sections then

treated with Gandy's method. In certain organs considerable increase of phosphatase activity was found. The most conspicuous increase in phosphatase activity could be observed in the liver which is negative otherwise. In addition phosphatase activity in the late capillaries and in the walls of the vessels. In other organs greater enzyme activity could be demonstrated in the basal membrane of the lungs in squamous in the transitional epithelium in the alveolar capillary blood and in desquamated alveolar cells of the lungs. Changes in activity were difficult to evaluate in other organs (atrophy of the stomach, difficulties in decalcification of bone marrow, great activity in the kidney of normal animals, etc.)

Conclusions: The increase in the phosphatase activity of the liver on irradiation can account for the hypoglycemic condition observed in connection with irradiations and the disappearance of liver glycogen in experimental animals. The increase in the phosphatase activity of the liver in the above mentioned conditions is connected with the direct action of the ionizing radiation and the secondary effects which presents itself in the form of a severe effect.

REFERENCES

H. Bauer 1946. 40. Thymus-Phosphatase. *Arch. für Vit.-Hormon-u. Enzymforsch.* 507-508. H.

T. Casparian, B. Thierl 1948. *Chromosoma* 13: 177. Quoted from H. Bauer Thymus-Phosphatase 568.

J. F. Danielli, D. G. Caspers 1945. *Nature*, 156, 298. Quoted from H. Bauer Thymus-Phosphatase 568.

F. Ellinger 1932. 181. *bid* Grundlagen der Strahlentherapie.

A. Klein, Ph. D. Kabat, J. Forth 1941. A histochemical study of the distribution of alkaline phosphatase in various normal and neoplastic tissues. *The Am. J. of Path.* 77: 303-318.

M. Ervov 1940. Action des rayons X sur la microchimie d'érythrocyte de poule. *Compt. rend. Soc. Biol.* 140.

M. Ervov, J. Brachet 1946. Action des rayons X sur la thymonucleotidase d'érythrocyte de poule in situ. *Compt. Rend. Soc. Biol.* 144.

R. Feudgen, H. Rosenfeld 1944. Mikroskopisch-chemischer Nachweis einer Nucleinase vom Typus der Thymonucleotidase und die damit verbundene selektive Färbung von Zellkernen in mikroskopischen Präparaten. *Zeits. für physiol. Chemie* 135: 201, 202.

S. J. Falley, H. D. Kay 1948. The phosphatases. *Enzymes* 4: 319-378. Verlag, Leipzig.

Gander 1939. Mikrotechnical demonstration of phosphatase in tissue sections. *Proc. Soc. Exp. Biol. & Med.* 42: 1.

Gander 1941. The distribution of phosphatase in normal organs and tissues. *J. of Cell Comp. Phys.* 17: 1.

Gander 1949. Histochemical significance of Phosphatases. *Proc. Soc. Exp. Biol. & Med.* 70: 1.

I. Hovoss 1945. *Rev. Med. Physiol.* XII: 102. Quoted from Dr. Lajtha I. A. *Enzymochromol. Gyvok Lappa* 1940, 1592. (Hung.)

B. Z. Hovoss 1947. The inhibition of ribo- and thymo-nucleic acid synthesis in tumor tissue by irradiation with X-rays. *Brit. J. Radiol.* 20: 438-450.

Dr. Lajtha I. 1936. A. *Enzymochromol. Gyvok Lappa* No. 30: 1509. (Hung.)

J. Rauber 1928. Über die Auswirkung der Röntgenstrahlenwirkung am biologischen Objekt. *Strahlentherapie* XXVII: 137, 236.

O. Strauss, J. Rauber 1924. Strahlentherapie auf das vegetative System. *Neurolog. Therapie* 18: 37.

ИЗМЕНЕНИЕ АКТИВНОСТИ ПЕЧЕНОЧНОГО ФАФАГАЗА В ОРГАНАХ КРЫС  
УБИТЫХ РАДИОНУКЛЕИДНЫМИ ЛУЧЕМАМИ

Докл. Докл. пр. Г. Рендерс, Вальтер  
Рейнхардт

В наших опытах мы исследовали крыс на их способность синтезировать  
нуклеинс. Мы подвергли крыс действию рентгеновских лучей количеством от 2000 до 4000  
Р поданных сразу или по частям (100 к В, 1 к А, 0,5 к С, 50 см). На каждое подопытное  
животное мы исследовали по одному контрольному животному. В части органов мы  
нашли значительное повышение активности ффафага. Исследуется также прежде всего  
проявление активности ффафага в печеночной ткани, где она обычно отсутствует, вли-  
же ограничивается на желчные капилляры и на стенку сосудов и в мышцах ло-  
желным следует повышение активности ффафага в печени под действием лучей  
по видимому объясняет биохимические состояния наблюдаемые под влиянием  
лучей и исчезновение активности в печени подопытных животных. Повышение активности  
ффафага в печени по видимому стоит в связи со детоксикацией йтимидилнуклеогенов,  
происходящей под влиянием лучей и связанной с действием йтимидина